information from Paul H. Moore, State FACE Project Officer, telephone (304) 285–6016, Internet: phm0@cdc.gov; Stephanie G. Pratt, State FACE Technical Officer, telephone (304) 285–5992, Internet: sgp2@cdc.gov, Trauma Investigations Section, Surveillance and Field Investigations Branch, Division of Safety Research, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention (CDC), 1095 Willowdale Road, Mailstop 180P, Morgantown, WV 26505–2888, or Nancy A. Stout, Ed.D., Director, telephone (304) 285–5894, Division of Safety Research, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention (CDC), 1095 Willowdale Road, Mailstop 1172, Morgantown, WV 26505–2888.

Potential applicants may obtain a copy of Healthy People 2000 (Full Report, Stock Number 017–001–00474–0) or Healthy People 2000 (Summary Report, Stock Number 017–001–00473–1) referenced in the **INTRODUCTION** through the Superintendent of Documents, Government Printing Office, Washington, DC 20402–9325, telephone (202) 512–1800.

Dated: January 5, 1998.

Diane Porter,

Acting Director, National Institute For Occupational Safety and Health Centers for Disease Control and Prevention (CDC).

[FR Doc. 98–3406 Filed 2–10–98; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97F–0522]

Anitox Corp.; Filing of Food Additive Petition (Animal Use)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Anitox Corp. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of formaldehyde in maintaining animal feeds and feed ingredients free of *Salmonella*.

DATES: Written comments on the petitioner's environmental assessment by March 13, 1998.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Henry E. Ekperigin, Center for Veterinary Medicine (HFV–222), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0174.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 2237) has been filed by Anitox Corp., P.O. Box 1929, Buford, GA 30519. The petition proposes to amend the food additive regulations in § 573.460 *Formaldehyde* (21 CFR 573.460) to provide for the safe use of formaldehyde (37 percent aqueous solution) at a maximum of 5.4 pounds per ton of animal feed and feed ingredients to maintain the animal feeds and feed ingredients free of *Salmonella*.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations promulgated under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (address above) for public review and comment. Interested persons may, on or before March 13, 1998, submit to the Dockets Management Branch (address above) written comments. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: January 26, 1998.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 98–3379 Filed 2–10–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98F–0063]

Protein Technologies International; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Protein Technologies International has filed a petition proposing that the food additive regulations be amended to provide for the safe use of a dry form of natamycin for use as an antimycotic in food.

FOR FURTHER INFORMATION CONTACT: JoAnn Ziyad, Center for Food Safety and Applied Nutrition (HFS–206), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3116.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 8A4581) has been filed by Protein Technologies International, Checkerboard Sq., St. Louis, MO 63164. The petition proposes to amend the food additive regulations in § 172.155 *Natamycin (pimaricin)* (21 CFR 172.155) to provide for the safe use of a dry form of the food additive for use on the surfaces of cuts and slices of cheese to inhibit mold spoilage, in accordance with various standards of identity for cheeses that allow the use of antimycotics and anticaking agents.

The agency has determined under 21 CFR 25.32(k) that this action is of the type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: January 26, 1998.

Laura M. Tarantino,

Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 98–3378 Filed 2–10–98; 8:45 am]

BILLING CODE 4160–01–F